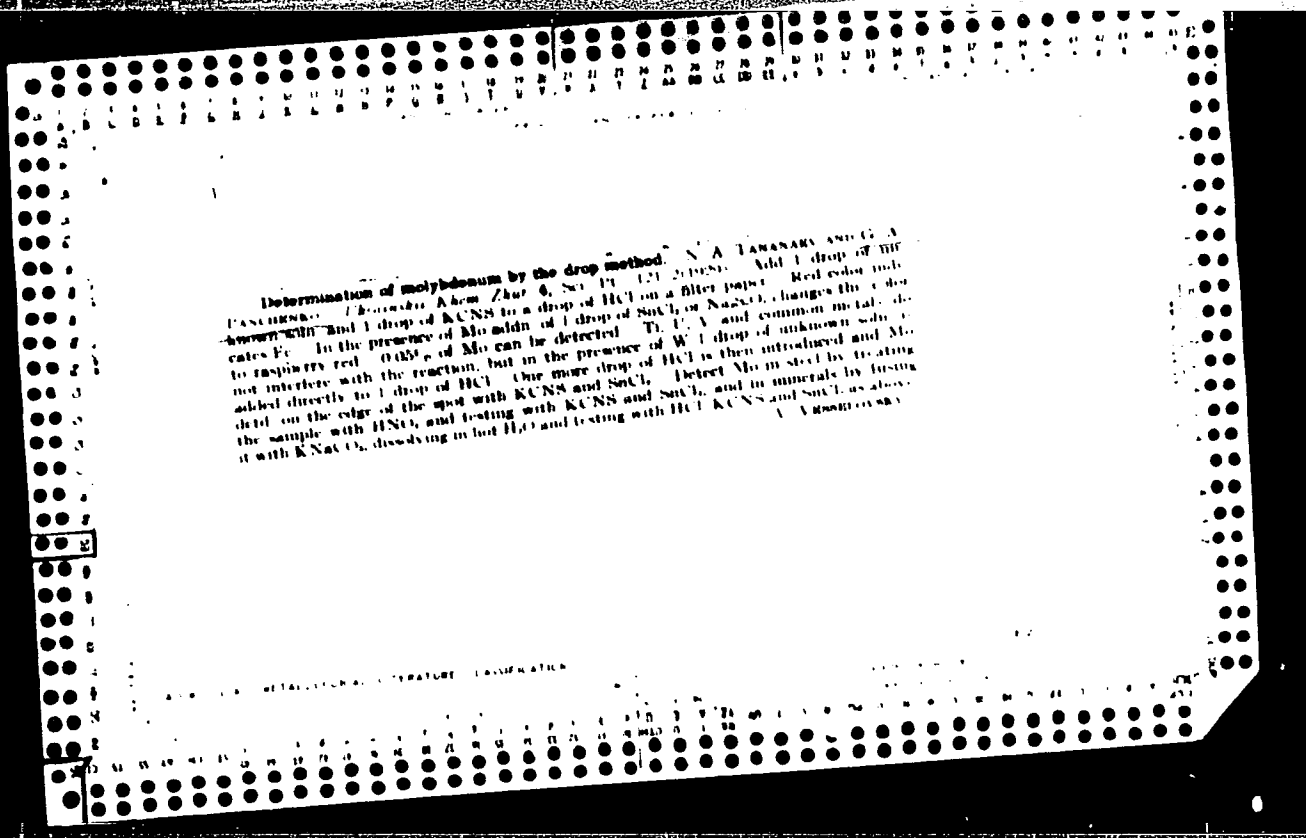




Detection of vanadium and tungsten. N. A. TANANAY AND G. A. PANCHENKO. *Russ. Pat. Chem. Soc.* 81, 1031-4 (1920). The most sensitive reaction for the detection of V takes place as follows. Take a strip of filter paper, introduce a drop of a mixt. of equal vols. of aniline and concd. HCl, then place over it a drop of the soln. to be tested which has previously been oxidized by boiling with concd. HNO₃ to make sure that V is not in the form of lower oxides, but in that of V₂O₅. The latter oxidizes aniline with the formation of an intense blue coloration. The sensitivity of this reaction is about 0.001 mg. metallic V per 1 drop (or 0.01 cc.) of the soln. Cations do not interfere with the reaction, but strong oxidizing anions, such as CrO₄, ClO₃, ClO₂, MnO₄, do interfere and must, therefore, be reduced by ebullition with HCl. HNO₃ does not interfere. To detect V in alloys, particularly in steels, a min. piece of the metal of the size of a pinhead is dissolved on a watchglass in aqua regia, the soln. is evaporated with an excess of HNO₃ to remove HCl, and one proceeds as above. Cr present in the alloy does not become transformed into CrO₃ and, therefore, does not interfere. When operating with minerals, the latter are dissolved either in aqua regia, if they contain little SnO₂, or by fusing with a mixt. of KNaCO₃ and KNO₃ and extracting with hot water, after which one may proceed as above, but in this case, if Cr was present, the chromate formed must be reduced by boiling with thiosulfate and HCl, after which V must be oxidized to V₂O₅ by boiling with HNO₃. To detect W in salts of H₂WO₄, place a drop of the soln. on a strip of filter paper, then place over it a drop of the soln. to be tested, a yellow spot is formed which, on being treated with SnCl₄, becomes blue colored. On adding NH₄CN, the blue coloration is intensified, and if it is, after that, treated with concd. HCl, it deepens still further. The sensitivity of this reaction is about 0.001 mg. of metallic W per drop (0.01 cc.) of the soln. If the soln. to be tested contains Mo, the action of SnCl₄ and NH₄CN causes the production of an intense cherry red coloration, but the latter disappears when concd. HCl is added, and thus does not interfere with the detection of W. To detect W in alloys, the latter are dissolved in aqua regia, the soln. is evaporated to dryness, the residue is calcined with osmic acid and, after being dissolved with water and NaOH, treated as above. W minerals are fused with KNaCO₃ and extracted with water, after which W can be detected as above. B. N.

PROCESSING AND PROPERTY DATA

Colorimetric determination of titanium in cast iron and steel. G. A. Pauzhenko and M. A. Rastvorov. *J. Appl. Chem.* (U.S.S.R.) 218, 21 (1955). Dissolve 10 ml of soln. (previously reduced with Hg/Zn to 30 ml), add 2 ml of 2% aq. chromotropic acid and compare the color with that given by 0.001 N Ti soln. The method involves an error of 0.03% for concns. of 5-20 mg of Ti per 100 ml. R. C. A.

ASAP-51A METALLURGICAL LITERATURE CLASSIFICATION

DATE: 11/11/55

RESEARCHER: R. C. A.

REVIEWER: R. C. A.

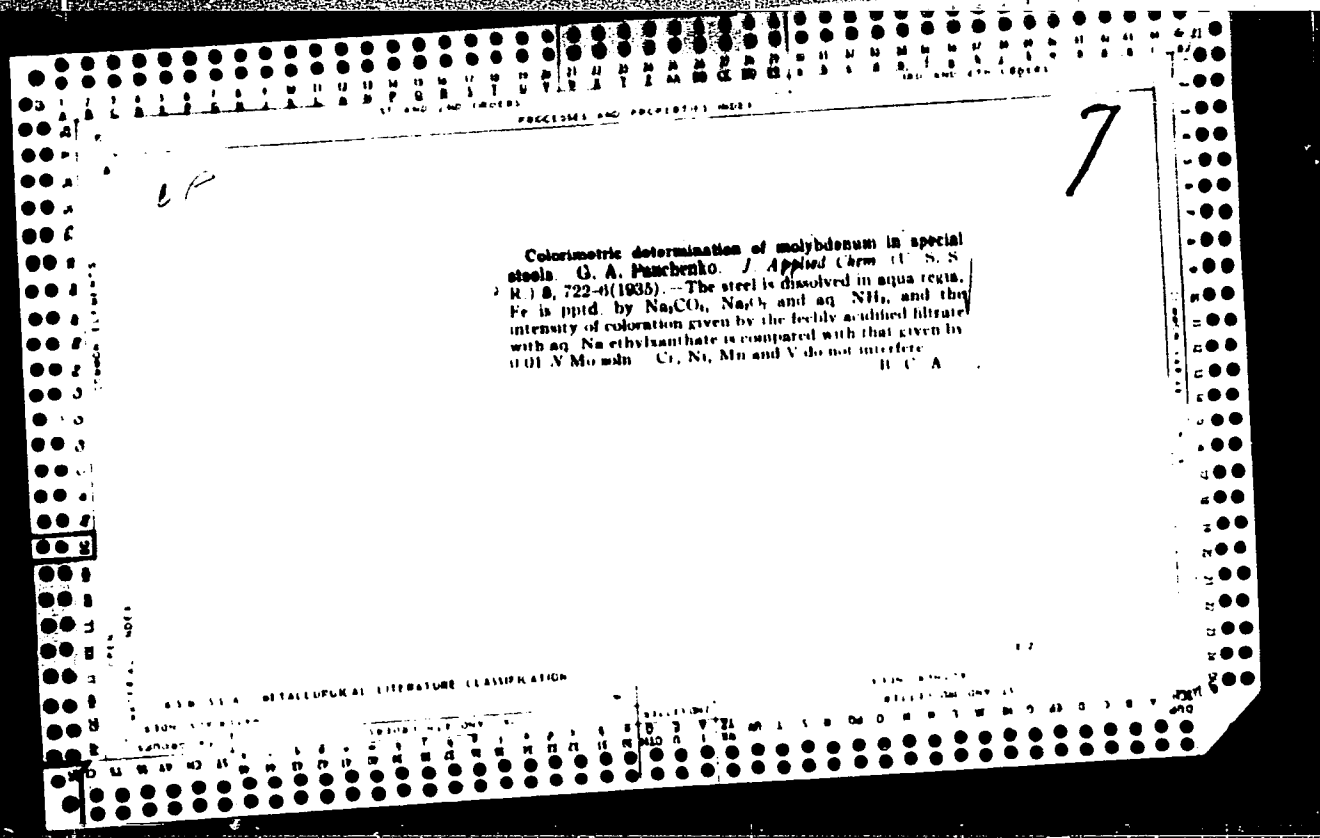
APPROVED: R. C. A.

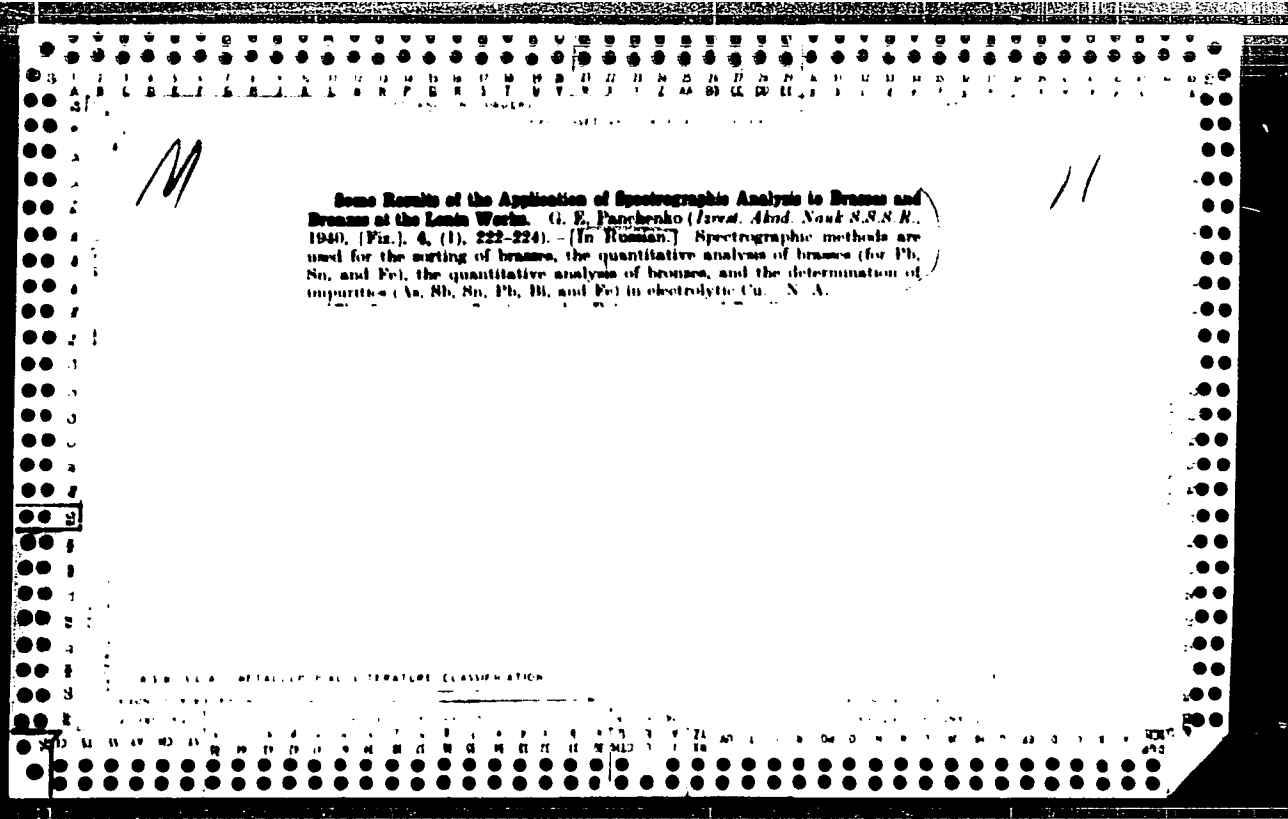
RECEIVED: R. C. A.

11

Isotometic determination of titanium, zirconium, niobium, hafnium, and vanadium in the presence of iron by means of liquid amalgams. A. A. Finklenberg, *J. Appl. Chem.* 1954, 6, 5, 535-536. (Received 1954, 10, 14). Methods for the determination of S, Se, Te, Pb, Bi, Al, and V in the presence of Fe, Cu, Ni, and Co in the form of liquid amalgams are described. The method is based on the formation of the amalgams of the elements to be determined with Fe, Cu, Ni, and Co. The amalgams are then dissolved in a suitable acid, and the solution is analyzed by means of a colorimetric method. The method is applicable to the determination of S, Se, Te, Pb, Bi, Al, and V in the presence of Fe, Cu, Ni, and Co in the form of liquid amalgams.

ALL INFORMATION CONTAINED HEREIN IS UNCLASSIFIED





1ST AND 2ND ORDERS

PROCESSES AND PROPERTIES

7

CH

Spectroscopy in the analysis of brasses and bronzes at the Lenin plant. G. R. Panchenko. *Dokl. akad. sci. U. R. S. S. Ser. phys.* 4, 222-4 (1940).—Spectral analysis is employed for sorting brasses and steels, according to trade marks, as well as for the analysis of brasses and electrolytic copper for impurities. Sorting of brasses is based mostly on detg. Cu and Zn, by making use of the lines Cu $\lambda = 3063 \text{ \AA}$. and Zn $\lambda = 3072 \text{ \AA}$. An activated λ arc is particularly suitable for the analysis. Rukhsana Gamow

COMMON ELEMENTS

MATERIALS

ASB-SLA METALLURGICAL LITERATURE CLASSIFICATION

BRONZES

STEELS

IRON

ALUMINUM

COPPER

ZINC

NICKEL

LEAD

TIN

SILVER

GOLD

PLATINUM

PALLADIUM

RHODIUM

IRIDIUM

OSMIUM

COBALT

NIOBIUM

TUNGSTEN

MOLYBDENUM

CHROMIUM

MANGANESE

IRON

STEELS

BRONZES

ALUMINUM

COPPER

ZINC

NICKEL

LEAD

TIN

SILVER

GOLD

PLATINUM

PALLADIUM

RHODIUM

IRIDIUM

OSMIUM

COBALT

NIOBIUM

TUNGSTEN

MOLYBDENUM

CHROMIUM

MANGANESE

Quantitative Spectrographic Analysis of Silicon Brasses for Silicon, Lead, and Zinc. (I. E. Panchenko (*Izvest. Akad. Nauk S.S.S.R.*, 1961, (Vis.), 5, (2,3), 313-318).—[In Russian.] A description is given of experiments carried out to determine suitable line pairs for use in the analysis of brasses containing Si 1-3-4, Pb 0-5-4, Zn 12-30, Fe 0-1-0-5%. Curves are given showing the time variation of these pairs. The Feussner spark was used as light source.—N. B. V.

FANCHENKO, G. E.

"Some Results of the Use of Spectroscopy in the

Analysis of Brasses and Bronzes at the Lenin Plant,"

Iz. Ak. Nauk SSSR, Ser. Fiz. 4, No. 1, 1940.

SIN'KOV, V.M., kand.tekhn.nauk; OKSANICH, M.A., inzh.; PANCHENKO, G.F., inzh.

Measuring relative increases of fuel consumption and efficiency of
boiler units. Avtom.i prib. no.2:113-103 '61. (MIRA 14:12)
(Boilers)

PANCHENKO, G.F.

Introducing a valveless proportioning pump. Biol. tekhn. zhurn. 1957.
Gos.nauch.-issl.inst.nauch.i tekhn.inform. 18 no.5:17-18 My 1956.
(MIRA 18:4)

PANCHENKO, G.F.

Unit for automatic control of automotive transportation in open
pi's. Biul.tekh.-ekon.inform.Gos.nauch.-issl.inst.nauch.1 tekhn.
inform. 18 no.1:65-66 Ja '65. (MIRA 18:4)

PANCHENKO, G.M., elektromekhanik; DEKHTENKO, Yu.K., elektromekhanik

A protective arc for telegraph apparatus. Avtom., telem. i
sviaz' 6 no.8:39 Ag '62. (MIRA 15:3)

1. Khar'kovskaya distantziya signalizatsii i svyazi Yuzhnoy dorogi.
(Telegraph--Equipment and supplies)

GUTNIKOV, B.Z., prof.; RUSAKOV, V.I., kand.med.nauk; PANCHENKO, G.S.,
kand.med.nauk; KOVALEV, G.G.; AKSEZHOV, A.I.; KHOPRYACHEV, N.V.;
KOMBACHEV, A.Sh.

Late results of treating patients with urethral strictures.
Urologia no.6:45-51 '60. (MIRA 15:5)

1. Iz fakul'tetskoy khirurgicheskoy kliniki (zav. - prof. B.Z.
Gutnikov) Rostovskogo meditsinskogo instituta.
(URETHRA--STRICTURE)

CHATSKIY, Ya.A.; PANCHENKO, G.T.

Surgical treatment of lung hemorrhage in pulmonary tuberculosis.
Probl.tub. no.4.106-107 1961. (MIRA 14.12)

1. Iz legochnogo otdeleniya (zaveduyushchiy Ya.A. Chatskiy)
Vinnitskogo oblastnogo tuberkuleznogo dispansera (glavnyy vrach
O.Z. Toretskaya).
(TUBERCULOSIS) (HEMORRHAGE) (LUNGS--SURGERY)

PANCHENKO, I.

Community workers check up. Okhr. truda i sots. strakh. 4 no.6:7-8
Je '61. (MIRA 14:7)

1. Predsedatel' zavkoma Lipetskogo traktornogo zavoda, g.
Lipetsk.

(Lipetsk—Insurance, Social)

(Lipetsk—Tractor industry—Hygienic aspects)

PANCHENKO, I.A.

Sensitivity of the respiratory center to carbon dioxide in the
state of chronic hypercapnia. Biul. eksp. biol. i med. 49
no. 5:25-28 My '60. (MIRA 13:12)

1. Iz fiziologicheskoy laboratorii (zav. - prof. L.L. Shik)
TSentral'nogo nauchno-issledovatel'skogo instituta eksperimentov
trudospособnosti i organizatsii truda invalidov (dir. - prof.
D.I. Gritskevich), Moskva. Predstavlena deystvitel'ny
chlenom AMN SSSR V.N. Chernigovskim.

(RESPIRATION) (CARBON DIOXIDE—PHYSIOLOGICAL EFFECT)

PANCHENKO, I.A.

Characteristics of respiration regulation after some surgical interventions on the chest. (Electromyographic study). Biul. eksp. biol. i med. 52 no. 11:27-30 N '61. (MIRA 15:3)

1. Iz fiziologicheskoy laboratorii (zav. - doktor meditsinskikh nauk Ye.N. Domontovich) Tsentral'nogo nauchno-issledovatel'skogo instituta ekspertizy trudosposobnosti i organizatsii truda invalidov (dir. - prof. D.I. Gritskevich), Moskva. Predstavlena deystvitel'nym chlenom AMN SSSR V.V. Parinym.
(RESPIRATION) (CHEST—SURGERY)
(ELECTROMYOGRAPHY)

PANCHENKO, I. D.

PA 64/49TB

USSR/Chemistry - Polarography Nov 48
Chemistry - Electrodes, Polarographic

"Use of Hard Electrodes in Polarography," Ye. M. Skobets, I. D. Panchenko, V. D. Ryabokon', Inst of Gen and Inorg Chem, Acad Sci Ukrainian SSR, 6 pp

"Zavod Lab" Vol XIV, No 11

Describes construction of a straight movable hard electrode manufactured either from silver or platinum amalgam. Determined that appearance of a maximum on the voltage curve which is characteristic for the mercury electrode is noticed only in a few cases with respect to the hard electrode.

64/49TB

PANCHENKO, I.D.

Use of solid electrodes for polarographic study of complex ions. Ukrain.
Khim. Zhur. 17, 872-6 '51. (MLRA 6:4)
(CA 47 no.22:12039 '53)

1. Inst. Gen. Inorg. Chem., Acad. Sci. Ukr. S.S.R., Kiev.

PANCHENKO, I. D.

USSR/Chemistry - Polarography

1 Jul 53

"The Applicability of the Heyrovsky-Il'kovich Equation to Polarographic Waves Taken at Solid Electrodes in Fused Salts," Yu. K. Del'marskiy, I. D. Panchenko, Inst of Gen and Inorg Chem, Acad Sci USSR

DAN SSSR, Vol 91, No 1, pp 115-118

Results obtained with the use of Pt electrodes on AgNO_3 , $\text{Cd}(\text{NO}_3)_2$, $\text{Mg}(\text{NO}_3)_2$, $\text{Zn}(\text{NO}_3)_2$, AgCl , CaCl_2 , TiCl_4 , PbCl_2 , ZnCl_2 , NiCl_2 , CoCl_2 , and CuCl_2

266T5

dissolved in molten NaNO_3 showed that the Heyrovsky-Il'kovich is valid for fused salts at solid electrodes. This opens up new possibilities for the polarographic investigation of salt melts. Presented by Acad A. N. Frumkin 24 Apr 53.

PANCHENKO, I. D.

Dissertation: "Polarographic Investigation of Molten Salts by Using Solid Electrodes."
Cand Chem Sci, Inst of General and Inorganic Chemistry, Acad Sci Ukrainian SSR, Kiev,
1953. (Referativnyy Zhurnal—Khimiya, Moscow, No 9, May 54)

SO: SUM 318, 23 Dec 1954

Panchenko, I. D.

V Diagrams of phase transformations of $RbCl-MgCl_2$ and $CsCl-MgCl_2$. B. P. Markov and I. D. Panchenko (Inst. Chem. and Inorg. Chem. Acad. Sci. USSR, Moscow, USSR). *Khim. Zhur.* 20, 620-3 (1964) (in Russian). -- These diagrams studied by visual-polythermic method show that 3 congruently melting compds., $MgCl_2 \cdot RbCl$ and $MgCl_2 \cdot 2RbCl$, are formed in the $RbCl-MgCl_2$ system; 3 compds., $MgCl_2 \cdot CsCl$, $MgCl_2 \cdot 3CsCl$, and $MgCl_2 \cdot 6CsCl$ and a congruently melting $CsCl \cdot 3MgCl_2$ are formed in the $CsCl-MgCl_2$ system. A definite regularity in diagram changes is observed when passing from $KCl-MgCl_2$ to $RbCl-MgCl_2$ and $CsCl-MgCl_2$; the crystn. of compds. extends over a greater range of compn., the crystn. temp. of compds. gradually increasing (cf. Menge, *C.A.*, 6, 44). Elizabeth Barabash

① 5/17

PANCHENKO, I. D.

USSR/ Chemistry - Physical chemistry

Card 1/1 : Pub. 147 - 16/22

Authors : Markov, B. F.; Delimarskiy, Yu. K.; and Panchenko, I. D.

Title : Thermodynamic properties of $PbCl_2$ in $PbCl_2$ - $LiCl$, $PbCl_2$ - $NaCl$, $PbCl_2$ - KCl , $PbCl_2$ - $RbCl$ -fusions.

Periodical : Zhur. fiz. khim. 28/11, 1987-1998, November 1954

Abstract : The electromotive forces of chemical chains with mixed electrolytes were measured in relation to temperature and composition of several binary lead chloride and alkali metal fusions. The thermodynamic properties of $PbCl_2$ in solutions with alkali metal chlorides were calculated. It was established, on the basis of thermodynamic data, that $PbCl_2$ - $LiCl$ solutions are almost ideal mixtures and that the components forming the solution blend together with the absorption of heat. The free reaction energy of $PbCl_2$ with alkali metal chlorides was determined. Eighteen references: 9-USSR; 6-German and 3-USA (1906-1953). Tables; graphs; drawing.

Institution : Academy of Sciences Ukr-SSR, Institute of General and Inorganic Chemistry

Submitted : March 21, 1954

MARKOV, B.F.; PANCHENKO, I.D.

Equilibrium diagrams of binary systems: magnesium chloride --
alkali metal chlorides. Zhur. ob. khim. 25 no.11:2038-2043 0
'55. (MLRA 9:4)

1. Institut obshchey i neorganicheskoy khimii Akademii nauk Ukrain-
skoy SSR.
(Chlorides)

PANCHENKO, I. D.

USER/ Chemistry - Physical chemistry

Card 1/2 Pub. 147 - 7/26

Authors : Markov, B. F.; Delimarskiy, Yu. K.; and Panchenko, I. D.

Title : Thermodynamic properties of $MgCl_2$ in $MgCl_2$ - $LiCl$, $MgCl_2$ - $NaCl$, $MgCl_2$ - KCl and $MgCl_2$ - $RbCl$ fusions.

Periodical : Zhur. fiz. khim. 29/1, 51-61, Jan 1955

Abstract : The electromotive forces of chemical chains with mixed $Mg/MgCl_2$ electrolytes were measured for various binary liquid systems and the thermodynamic properties of $MgCl_2$ were calculated in solutions with alkali metal chlorides. It was found that $MgCl_2$ and $LiCl$ create solutions close to ideal mixtures. Data are given on the partial isobaric potential of $MgCl_2$ as well as its partial entropy.

Institution : Academy of Sciences Ukr SSR, Institute of General and Inorg. Chem., Kiev.

Submitted : March 20, 1954

Periodical : Zhur. fiz. khim. 29/1, 51-61, Jan 1955

Card 2/2 Pub. 147 - 7/26

Abstract : The thermodynamic properties of $MgCl_2$ in solutions with KCl and $RbCl$ indicated a deep reaction between the individual components which led to the formation of compounds capable of being separated in the solid state. Eight references; 4 USSR; 3 German and 1 Swiss. (1911-1954). Tables; graphs; drawing.

Panchenko, I. D.

USSR/Physical Chemistry - Electrochemistry, B-12

Abst Journal: Referat Zhur - Khimiya, No 1, 1957, 536

Author: Panchenko, I. D.

Institution: None

Title: On the Equation of the Polarographic Wave Obtained with a Hard Electrode in Fused Salts

Original

Periodical: Ukr. khim. zh., 1956, Vol 22, No 2, 153-155

Abstract: On the basis of the picture of the deposition of metals from fused salts during polarographic recording with a hard electrode presented earlier (Referat Zhur - Khimiya, 1953, 2881; 1954, 21433; 1955, 5357), a more exact equation for the polarographic wave under these conditions is given. The equation is based on the assumption that the activity of the deposited metal is a function of the current, due to the diffusion of the metal in the hard electrode. The equation thus obtained is identical with the Geyrov-Il'kovich equation.

Card 1/1 *Inst. obshey. neorganicheskoy khimii AN USSR.*

PANCHENKO, J. D.

1. The first part of the document is a list of names of persons who were in contact with the subject of the investigation. The names are listed in alphabetical order. The names are: [illegible]

PANCHENKO, I. D., MARKOV, B. F., and DELIMARSKIY, Y. K., Institute of General
and Inorganic Chemistry, AS USSR, Kiev.

"Thermodynamic Properties of Certain Chlorides in Melts,"
a paper submitted at the International Symposium on Macromolecular
Chemistry, 9-15 S p 1957, Prague.

MARKOV, B. G., DELIMARSKIY, U. K., and FANCHENKO, I. D.

(Institute of General and Inorganic Chemistry, Acad. Sci. Ukr. SSR, Kiev, USSR)

"Thermodynamic Properties of Some Fused Chlorides,"
paper submitted at Soviet High-Polymers, Intl. Conference, Nottingham,
21-24 July 1958.

E-3,109,661

PANCHENKO, I.D.

Investigating the performance of lubrication system valves in
D-54 engines. Mekh. sil'. nos. 9 no.4:26-27 Ap '58. (MIRA 11:5)

1. Ukrains'ka akademiya sil'skogospodarskikh nauk
(Tractors--Eng⁴nes)

I. D. PANCHENKO, GITMAN, Ye. B., A. A. KOLOTTIY, Yu. K. DELIMARSKIY

"Electrolytic Production of Lead by Electrolytes of Fused Salts"

IONKh Ac. Sc. Ukr SSR.

report submitted at a conference on new methods of lead production from concentrates,
Gintsvetmet (State Inst. Non-Ferrous Metallurgy), Moscow 22-25 June 1958.

(for entire conf. see card for LIDOV, V. P.)

PHASE I BOOK EXPLOITATION SOV/2.17
 PANCHENKO, G. P.

5(4)
 PHASE I BOOK EXPLOITATION SOV/2.17
 Soveshchaniye po elektrokhemii. 4th, Moscow, 1956.
 Trudy... [sbornik] (Transactions of the Fourth Conference on Electrochemistry: Collection of Articles) Moscow, Izd-vo AN SSSR, 1959. 868 p. Errata slip inserted. 2,500 copies printed.
 Sponsoring Agency: Akademiya nauk SSSR. Otdeleniye khimicheskikh nauk.

Editorial Board: A. M. Prumkin (Resp. Ed.) Academician, O. A. Yasin, Professor, S. I. Zhdanov (Resp. Secretary), B. M. Kabanov, Professor, S. I. Zhurav (Resp. Secretary), D. M. Kabanov, Professor, Ya. M. Kolotyrkin, Doctor of Chemical Sciences, V. V. Loshakov, P. D. Lukovtsev, Professor, Z. A. Solov'yeva, V. V. Stender, Professor, and G. M. Florinovich; Ed. of Publishing House: N. G. Yegorov; Tech. Ed.: T. A. Prusakova.

PURPOSE: This book is intended for chemical and electrical engineers, physicists, metallurgists and researchers interested in various aspects of electrochemistry.

COVERAGE: The book contains 127 of the 138 reports presented at the Fourth Conference on Electrochemistry sponsored by the Department of Chemical Sciences and the Institute of Physical Chemistry and Academy of Sciences, USSR. The collection pertains to different branches of electrochemical kinetics, double layer theories and galvanic processes in metal electrodes, the end of each division. Abridged discussions are given at the end of each division. The majority of reports not included here have been published in periodical literature. No personalities are mentioned. References are given at the end of most of the articles.

Golubev, Ya. P. (Institut geokhimi i analiticheskoy khimii AN SSSR, imeni V. I. Vernadskogo, Institute of Geochemistry and Analytical Chemistry, imeni V. I. Vernadskiy, Academy of Sciences, USSR). Diffusion of Electrolytes and the Polarographic Method 677

Rosentzweig, T. and K. A. Zhigalova (Institute of Physical Chemistry, Academy of Sciences, USSR). Diffusion of Gases Through Thin Films of Electrolytes 684

Discussion 10.8. Ksenzhuk, Yu. A. Chismadshay, Yu. A. Vdovin, O. B. Khachaturyan and contributing authors 689

PART VIII. ELECTROCHEMICAL PROCESSES IN NONFERROUS METALLURGY 695

Stender, V. V. (Dnepropetrovsk Institute of Chemical Technology imeni P. E. Dzerzhinskogo, Institute of Chemistry, Academy of Sciences, KazSSR). Electrolysis as a Means of Combining Card 27/34

Several Metallurgical and Chemical Production Processes (Some New Processes of Hydroelectric Metallurgy) 697

Kozlovskiy, M. T. (Kazakh State University, Academy of Sciences, KazSSR). Some Problems of Amalgam Metallurgy: Cementation of Metals With Amalgam 704

Dellmar, Yu. M., B. P. Markov, I. D. Panchenko, Ye. B. Giman, and A. A. Agabekov (Institute of General and Organic Chemistry, Academy of Sciences, KazSSR). Electrolytic Purification of Lead from Paved Sulfate 710

Chishikov, D. M. and V. M. Koylina (Institute of Metallurgy, Academy of Sciences, USSR). Investigation of the Potentials and Anodic Polarization of Metallic Sulfides and Their Alloys 715

Lesin, T. I. and I. A. Bauman (Deceased) (Vsesoyuznyy nauchno-issledovatel'skiy institut tvoricheskoy metallurgii, All-Union Scientific Research Institute of Nonferrous Metals, Sverdlovsk) Card 28/34

Features of the Anode Process During the Purification of a Copper-Nickel Anode in a Sulfate-Chloride Electrolyte 720
 Zaretskiy, S. A., I. G. Zharul'skiy (Deceased), and I. A. Bogdanova. Anodic Behavior of Manganese and Its Alloys 724

PANCHENKO, I. D.

5(4) PHASE I BOOK EXPLOITATION SUB

Soveshchaniye po elektrokhemii. 4th, Moscow, 1956.

Trudy... [Izbornik] (Transactions of the Fourth Conference on Electrochemistry: Collection of Articles) Moscow, Izd-vo AN SSSR, 1959. 868 p. Errata slip inserted. 2,500 copies printed. Sponsoring Agency: Akademiya nauk SSSR. Otdeleniye khimicheskoy nauk.

Editorial Board: A.M. Prumkin (Resp. Ed.) Academician, O.A. Yasin, Professor, S.I. Zhdanov (Resp. Secretary), B.M. Kabanov, Professor, S.I. Zhdanov (Resp. Secretary), B.M. Kabanov, Professor, Ya. M. Kolotyrkin, Doctor of Chemical Sciences, V.V. Jonev, P.D. Lukovitsky, Professor, Z.A. Solov'yeva, V. Standart, Professor, and O.M. Florianovich, Ed. of Publishing House M.O. Yegorovi; Tech. Ed.: T.A. Prusakova.

PURPOSE: This book is intended for chemical and electrical engineers, physicists, metallurgists and researchers interested in various aspects of electrochemistry.

COVERAGE: The book contains 127 of the 138 reports presented at the Fourth Conference on Electrochemistry sponsored by the Department of Chemical Sciences and the Institute of Physical Chemistry and Academy of Sciences, USSR. The collection contains to different branches of electrochemical kinetics, double layer theories and galvanic processes in metal electrodes, and industrial electrolysis. The majority of reports are included here have been published in periodical literature. No personalities are mentioned. References are given at the end of most of the articles.

Golubtsev, Ya. P. (Institut geokhimi i analiticheskoy khimii AN SSSR, Leningrad), V.I. Vernadskogo - Institute of Geochemistry and Analytical Chemistry, Leningrad, V.I. Vernadskiy, Academy of Sciences, USSR). Diffusion of Electrolytes and the Polarographic Method 677

Rogachev, I. T. and K.A. Zhizalova (Institute of Physical Chemistry, Academy of Sciences, USSR). Diffusion of Oxygen Through Thin Films of Electrolytes 684

Discussion (O.S. Ksenzhek, Yu. A. Chizmadzhev, Yu. A. Vdovin, O.B. Khachaturyan and contributing authors) 689

PART VIII. ELECTROCHEMICAL PROCESSES IN NONFERROUS METALLURGY 695

Stender, V. V. (Dnepropetrovsk Institute of Chemical Technology - Iseni P. E. Dzerzhinskii, Institute of Chemistry, Academy of Sciences, KazSSR). Electrolysis as a Means of Combining 697

Card 27/34

Several Metallurgical and Chemical Production Processes (Some New Processes of Hydroelectric Metallurgy) 697

Mozlovskiy, M. T. (Kazakh State University, Academy of Sciences, KazSSR). Some Problems of Amalgam Metallurgy - Cementation of Metals With Amalgams 704

Delmarakiz, Yu. K., B. P. Markov, I. D. Panchenko, Ye. B. Dilman, and A. A. Ryabitskiy (Institute of Chemistry, Academy of Sciences, KazSSR). Electrolytic Purification of Lead from Paved Salts 710

Chizhikov, D. M., and V. N. Koryulina (Institute of Metallurgy, Academy of Sciences, USSR). Investigation of the Potentials and Anodic Polarization of Metallic Sulfides and Their Alloys 715

Lesin, P. I., and I. A. Buzman (Deceased) (Vsesoyuznyy nauchno-issledovatel'skiy institut tsvalykh metallov, All-Union Scientific Research Institute of Nonferrous Metals) Special 715

Card 28/34

Features of the Anodic Process During the Purification of a Copper-Nickel Anode in a Sulfate-Chloride Electrolyte Zaretakiy, S. A., I. G. Zhuritskiy (Deceased), and I. A. Bogdanova (Deceased) 721

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S/075/60/015/004/002/030/XX
B020/B064

26.620
AUTHOR:

Panchenko, I. D.

TITLE:

Derivative Polarography on Solid, Rotating Electrodes in
Molten Salts

PERIODICAL:

Zhurnal analiticheskoy khimii, 1960, Vol 15, No 4,
pp. 388 - 390

TEXT: Since a number of metals is obtained by electrolysis of molten salts, the investigation of the electrochemical properties of the accompanying substances is of interest, which is possible in each case by means of derivative polarography. Ye. M. Skobets (Ref. 1) obtained the derivation peaks for a number of compounds on solid electrodes and in aqueous solutions, and N. G. Chovnyk (Ref. 2) in molten salts. The author aimed at determining the decomposition voltage of some metal chlorides, studying the change of the diffusion current as dependent on the speed of rotation of the electrodes, and the change of the polarization curves as dependent on the rate of polarization. Silver, lead, and cadmium chlorides were investigated; a eutectic LiCl-KCl mixture was used as

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supporting electrolyte, and the polarograms recorded with a Heyrovský polarograph. The sensitivity of the galvanometer was $1.6 \cdot 10^{-9}$ a/mm. The investigations were conducted at 400°C , and the temperature was kept constant to within $\pm 2^\circ\text{C}$ by means of a thermostat. Porcelain crucibles with a platinum wire cathode and a platinum plate anode served as electrolyzers. The electrodes had a speed of 100, 500, 750, and 1500 rpm. Fig. 1 gives the scheme of the polarographing cell. The authors first recorded the usual polarograms with separate and joint precipitation of the metals on an LiCl-KCl background. With separate precipitation of the metals, reproducible polarograms were obtained, whereas with joint precipitation no positive results were obtained. The decomposition voltage of the supporting electrolyte equals -2.3 v. The half-wave potential for silver is 0.9 v, and for lead, 0.95 v. In the reduction of cadmium chloride, two waves were obtained; the potential of the first half-wave was 1.1 v, and that of the second one, 1.3 v. In the authors' opinion, the first wave corresponds to the reduction of cadmium chloride to the subchloride, and the second to the reduction of the subchloride to metallic cadmium. At a concentration of 0.001 molar parts, the diffusion current limits were

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0.24 ma for silver, 0.32 ma for lead, and 0.16 and 0.06 ma, respectively, for cadmium. The derivative polarograms were recorded at a sensitivity of $S=1/200$. The derivation polarograms of silver precipitated from an LiCl-KCl supporting electrolyte at an AgCl concentration of 0.001 molar parts (Fig. 2) and of cadmium precipitated from an LiCl-KCl supporting electrolyte at a $CdCl_2$ concentration of 0.0007 molar parts (Fig. 3) are

given. The derivative peak potentials of the chlorides investigated correspond to those of the half-waves with insignificant deviations. The derivative peaks (Fig. 4) resulting from coprecipitation of silver and cadmium are similar to those found in coprecipitation of lead and cadmium. With a rotating cathode, the height of the derivative peak as compared to the stationary electrode is twice as much at 100 rpm, at 500-750 rpm, three times as much, and at 1500 rpm, four times as much. The respective derivative polarograms are shown in Fig. 5. An exact mathematical relation between the speed of the electrode and the height of the derivative polarogram peak cannot be established. The diffusion currents on rotating electrodes in aqueous solutions increase 15-20 times at 700-800 rpm; diffusion currents in melts are, therefore, 5-7 times smaller than in

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aqueous solutions, which is due to the temperature of the former since the diffusion coefficient is an exponential function of temperature. Since the coprecipitation of the metal ions examined showed no positive results on ordinary polarograms, the polarograms were recorded for low rates of polarization (10-20 mv/min), and it was found that, at low rates, the limiting currents are recorded in the coprecipitation of metals (Fig. 6). and the stepwise reduction of cadmium chloride is confirmed. This paper was read in the Analytical Section of the VIII Mendeleyevskiy s"yezd (VIII Mendeleyev Congress) on March 21, 1959. There are 6 figures and 8 references: 6 Soviet, 1 US, and 1 German.

ASSOCIATION: Institut obshchey i neorganicheskoy khimii AN USSR, Kiyev
(Institute of General and Inorganic Chemistry of the
AS UkrSSR, Kiyev)

SUBMITTED: June 15, 1959

Card 4/4

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77517
SCN 50-33-1-260-49

AUTHORS: Panchenko, I. D., Delimarskiy, Yu. K.

TITLE: Electrolytic Recovery of Lead From the Factory Crude Lead and Bismuth Dross Using Melted Electrolytes

PERIODICAL: Zhurnal prikladnoy khimii, 1960, Vol 33, Nr 1, pp 153-156 (USSR)

ABSTRACT: Large-scale laboratory experiments of electrolytic lead refining were conducted in order to find the best conditions and materials for removing impurities from lead and for producing different grades of lead. Crude factory lead and bismuth dross were used. The following conclusions were made from the data obtained. These eutectic electrolytes may be used in electrolytic lead refining: $PbCl_2$ -KCl-NaCl, $PbCl_2$ -NaCl, and $PbCl_2$ -PbO.

It was found that by refining the crude lead with 25% recovery of the cathode lead, grade C_1 lead is obtained; with 50% recovery, grade C_2 lead is obtained; with 75% recovery in a triple and double eutectic mixture of lead,

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Factory Crude Lead and Bismuth Dross
Using Melted Electrolytes

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potassium, and sodium chlorides, grade C_3 lead is obtained. It was found that by the electrolytic refining of bismuth dross containing 5% of Bi, grade C_1 lead can be obtained (one thousandth of a percent of Bi); dross containing 13% of Bi yields grade C_2 cathode lead (containing one hundredth of a percent of Bi). It was found that the following electrolysis conditions are optimal: temperature, 500°C ; anode current density (D_a), 0.4 amp/cm²; cathode current density (D_c), 1 amp/cm². Yield based on current, 97-98%. Electric energy consumption, 1.6 kw hours/kg. There are 4 tables; and 6 Soviet references.

SUBMITTED: November 15, 1958
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S/021/61/000/002/01.017
D210/D303

AUTHORS: Delimars'kyyi, Yu.K., Member of AS UkrSSR, Pachenko
I.D., and Shylina, H.V.

TITLE: Rotating disc electrode in the polarography of
fused salts

PERIODICAL: Akademiya nauk Ukrayins'koyi RSR. Dopovidi, no. 2
1961, 205 - 208

TEXT: The authors state that the question of theoretically interpreting polarographical curves obtained for fused electrolytes with the use of stationary or revolving cylindrical electrodes cannot be regarded as solved, because different investigators propose different equations for their interpretation. Only V.G. Levich (Ref. 3: Fiziko-khimicheskaya gidrodinamika, Fizmatgiz, M., 1960 (Physico-Chemical Hydrodynamics)) worked out a theory of convective diffusion of a rotating disc electrode, its equity having been proved for aqueous electrolyte solutions. The thickness of the

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diffusion layer is given by the equation

$$\delta = 1.6 \cdot D^{1/3} \cdot \gamma^{1/6} \cdot \omega^{1/2}$$

where δ is the layer thickness, D - coefficient of diffusion, γ - kinematic viscosity, ω - angular velocity of the rotating disc electrode. The ultimate current value is given by the equation

$$i_d = \frac{nFDC}{\delta}$$

which, taking into account Eq. (4) may be expressed as

$$i_d = 0.62 nFD^{2/3} \cdot \gamma^{-1/6} \cdot \omega^{1/2}$$

The authors decided to apply Levich's theory to fused salts. They obtained polarograms of a series of electrolytes, using a rotating disc electrode. They investigated AgCl , PbCl_2 , CdCl_2 and TiCl_4 on the background of a fused mixture of Cl_2 - KCl . Polarograms were

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obtained by means of a check polarograph LP55A. Experiments were carried out at 420°C in a china crucible, cap. 100 ml, the anode was a platinum plate of 4 cm², the cathode a platinum microelectrode set in a molybdenum tube, disc's angular velocity 100, 500 and 1000 rev/min. Apart from normal polarograms they obtained derivative ones, by using an electrolytic condenser, consecutively connected with the galvanometer. The depolarizer concentrations were in the range of 0.0005 to 0.003 mol fractions. Their first object was to investigate the dependence of the limit-current on the rate of electrode revolutions. Polarograms obtained with different electrode angular velocities are shown in Fig. 1. On the basis of these, the authors found that the above dependence can be expressed by the equation:

$$i_d = K \cdot \omega^{0.58}$$

This is in agreement with Eq. (6), which follows from Levich's connective diffusion theory. They thus proved the adequacy of this theory for fused electrolytes, in which it is possible to use a rotating disc electrode.

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tating disc electrode. Their experimental results are given in tabulated form. Two waves were observed for cadmium and lead chlorides which is obviously due to the formation of univalent chlorides. The authors express the opinion that the use of the rotating disc electrode allows polarographic curves to be obtained with lower concentrations than with the stationary ones which is a proof of its higher susceptibility and permits a more precise polarogram interpretation. Kinematic viscosity values being known, it is possible to calculate from Eq. (6) the diffusion coefficients in fused electrolytes; the authors found that this coefficient for silver ions equals $0.9 \cdot 10^{-5} \text{ cm}^2/\text{sec}$, which is in good agreement with data obtained for determining it by different methods. [Abstractor's note: Methods not given]. There are 3 figures, 1 table, and 8 references: 6 Soviet-bloc and 2 non-Soviet-bloc. The references to the English-language publications read as follows: E. D. Black, T. De-Vries, *Analyt. Chem.*, 27, 906, 1955; J. Corbett, S. Wintash, F. Alberts, *J. Am. Chem. Soc.*, 79, 3020, 1957.

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Rotating disc electrode in ...

S/021/61/000/002/011/013
D210/D303

ASSOCIATION: Institut zagal'noyi ta neorganichnoyi khimiyi AN URSR
(Institute of General and Anorganic Chemistry, AS
UkrSSR)

SUBMITTED: September 3, 1960

Card 5/6

PANCHENKO, I.D., kand. tekhn. nauk; VEDMID', M.P., kand. tekhn. nauk;
NATANZON, I.I., kand. tekhn. nauk, red.; KOSOVSKIY, V.A.
[Kosovs'kyi, V.A.], red.; KVITKA, S.P., tekhn. red.

[Temperature conditions of a lubrication system and their
regulation] Temperaturnyi rezhym systemy mashchennia ta ioho
reguliuвання. Kyiv, Vyd-vo Ukr. Akad. sil's'kohospodars'kykh
nauk, 1961. 146 p. (MIRA 15:4)
(Tractors--Lubrication)

PANCHENKO, I.D.; SHILINA, G.V.

Rotating disk electrode in the polarography of fused salts.
Zhur.anal.khim. 18 no.8:920-923. Ag '63. (MIRA 16:12)

1. Institute of General and Inorganic Chemistry, Academy of
Sciences, Ukrainian S.S.R., Kiev.

PANCHENKO, I.D.; SHILINA, G.V.

Theory of convective diffusion in the polarography of fused salts. Ukr. khim. zhur. 29 no.11:1164-1169 '63. (MIRA 16:12)

1. Institut obshchey i neorganicheskoy khimii AN UkrSSR.

L 30243-66 EWT(m)/ETC(f)/T/EMP(t)/ETI IJP(c) DS/JD/JG

ACC NR: AP6013884

SOURCE CODE: UR/0073/65/031/011/1203/1206

AUTHOR: Panchenko, I. D.; Penkalo, I. I.

ORG: Institute of General and Inorganic Chemistry, AN UkrSSR (Institut obshchey i neorganicheskoy khimii AN UkrSSR)

TITLE: Polarographic study of rare earth elements in fused potassium bisulfate as the supporting electrolyte

SOURCE: Ukrainskiy khimicheskiy zhurnal, v. 31, no. 11, 1965, 1203-1206

TOPIC TAGS: polarography, lanthanum oxide, gadolinium compound, yttrium compound, cerium compound, europium compound, potassium compound / LP-55A polarograph

ABSTRACT: A polarographic study of oxides of lanthanum, gadolinium, yttrium, cerium, and europium was carried out in potassium bisulfate as the supporting electrolyte by using a rotating disc electrode. The reduction polarograms were recorded on an LP-55A polarograph at 240°C, and the half-wave potentials of the reduction of the oxides were determined: $E_{1/2}$ was 1.1 V for La_2O_3 , 1.08 V for Gd_2O_3 , 1.03 V for Y_2O_3 , 0.9 V for

CeO_2 , and 0.4 V for Eu_2O_3 . Thus, the deposition potentials differed appreciably in this fused electrolyte. The limiting diffusion current was found to be directly proportional to the concentration of the oxides studied. The diffusion coefficients of

UDC: 543.253+546.6+541.133

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ACC NR: AP6013884

the ions were found to be $2.6 \cdot 10^{-5}$ for La, $0.31 \cdot 10^{-5}$ for Gd, $0.33 \cdot 10^{-5}$ for Y, $1.11 \cdot 10^{-5}$ for Ce, and $1.34 \cdot 10^{-5}$ for Eu. The thickness of the diffusion layer in fused potassium bisulfate was calculated to be $\sim 2 \cdot 10^{-3}$ cm. Orig. art. has: 5 figures, 1 table.

SUB CODE: 07/ SUBM DATE: 26Oct64/ ORIG REF: 006/ OTH REF: 002

Card 2/2 CC

PANCHENKO, I.D.; BOYKO, K.M.

Use of various electrode materials in the polarography of fused salts. Ukr.khim.zhur. 31 no.10-194 '65.

(MIRA 18:4)

1. Institut obshchey i neorganicheskoy khimii AN UkrSSR.

PANCHENKO, I. S.

Use of an automatic metal dropping electrode in the polarography
of fus. salts. Zhur. fiz. khim. 39 no.2:514-515 F '65.
(MIRA 18:4)

1, Institut issledovaniy v organicheskoy khimii AN SSSR.

L 30215-66 EWT(m)/ETC(f)/T/EWP(t)/ETI IJP(c) DS/JD/JG
 ACC NR: AP6015011 SOURCE CODE: UR/0364/66/002/005/0529/0535

AUTHOR: Panchenko, I. D.; Penkalo, I. I.; Delimarskiy, Yu. K.

ORG: Institute of General and Inorganic Chemistry, AN UkrSSR, Kiev (Institut obshchey i neorganicheskoy khimii AN UkrSSR)

TITLE: Polarographic study of the cerium group of rare earth elements in the fused LiCl-KCl eutectic as supporting electrolyte

SOURCE: Elektrokimiya, v. 2, no. 5, 1966, 529-535

TOPIC TAGS: polarographic analysis, lanthanum compound, praseodymium compound, samarium compound, europium compound, cerium compound, neodymium compound, lithium chloride, potassium chloride

ABSTRACT: In order to determine its usefulness for analytical purposes, the authors investigated the polarographic behavior of lanthanum, cerium, praseodymium, neodymium, samarium, and europium in the fused LiCl-KCl eutectic at 400°C, using a rotating platinum disc electrode. Polarograms of the reduction of the rare earth chlorides were recorded at various concentrations of the chlorides. A direct proportion was established between the wave height and the concentration of the depolarizer. The half-wave potentials were determined and differences between some of them were found to exceed

UDC: 541.135.3

Cord 1/2

SOV/149-58-5-14/18

AUTHORS: Layner, V.I., Panchenko, I.I.

TITLE: Electrode Processes in Electro-deposition of Nickel from Fluoborate Electrolytes (Elektroodnyye protsessy pri elektro-osazhdenii nikelya iz ftorboratnykh elektrolitov)

PERIODICAL: Izvestiya Vysshikh Uchebnykh Zavedeniy, Tsvetnaya Metallurgiya, 1958, Nr 5, pp 124 - 130 (USSR)

ABSTRACT: The object of the investigation described in this paper was to study the effect of various factors on electrodeposition of nickel from fluoborate solutions and on the quality of the deposits obtained by this method. For the preparation of the HBF_4 solution, chemically pure H_3BO_3 and an HF solution (whose concentration was determined from its density) were used in the stoichiometric ratio, H_3BO_3 being added (a small quantity at a time) to the continuously stirred and ice-cooled HF solution. To the obtained HBF_4 solution, also continuously cooled, nickel carbonate (a small quantity at a time) was added and in this manner it was possible to obtain solutions containing up to 180 g/litre nickel in the form of

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Electrode Processes in Electro-deposition of Nickel from Fluoborate Electrolytes

fluoborate (density approx. 1.5), which could then be diluted to any required concentration. The characteristics of the experimental electrolytes (pH = 3 in all cases) are given in Table 1 which shows the nickel concentration ($N = 1$ to 4), the content (in g/litre) of fluorine, F_1 , present in the form of BF_3OH , the content of fluorine, F_2 , present in the form of BF_4^- , the total fluorine content $F = F_1 + F_2$, the B content, the F/B and F/Ni ratios, the $NiCl_2 \cdot 6H_2O$ content, and the density at $20^\circ C$. Nickel anodes and steel, copper or brass cathodes were used in the experiments, the results of which are reproduced graphically. The effect of the current density (A/dm^2) on the cathode potential is illustrated in Figure 1, where graphs 1, 2, 3 and 4 correspond to the nickel concentrations of 1N, 2N, 3N and 4N, respectively. Figure 2 illustrates the current-density/cathode-potential relationship for electrolytes with no excess of H_3BO_3

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(graph 1) and with 5, 10 and 15g/litre H_3BO_3 in excess of the stoichiometric ratio (graphs 2,3 and 4). The results of the tests in which the current-density/cathode-potential relationship was studied for 3N electrolytes containing no Cl ions at 20 °C and those containing 15, 30 and 50 g/litre $NiCl_2 \cdot 6H_2O$ at 50 °C are reproduced in Figure 3 (graphs 1, 2, 3 and 4, respectively). The same relationship for a 3N electrolyte with no excess of H_3BO_3 at 20, 30, 40 and 50 °C is shown in Figure 4 (graphs 1 to 4). A 3N electrolyte was also used for investigating the effect of the pH number which was varied between 1 and 5 by means of HBF_4 or sodium iodide additions. It was found that at $pH^4 = 5$, the electrolyte is unstable and contains black, insoluble particles (most likely $Ni(OH)_2$ with basic salts) held in suspension. Pitting occurs and a dendritic deposit is obtained which at low current densities (1 to 5 A/dm^2) becomes dark. At $pH = 4$, low current

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Electrode Processes in Electro-deposition of Nickel From Fluoborate Electrolytes

densities result in a dark deposit, while dendrites are formed at high current densities. At $\text{pH} = 1$ or 2 , intensive evolution of hydrogen takes place on the cathode surface, the evolved gas forming bubbles which adhere to the cathode and cause pitting. Best quality deposits are obtained at $\text{pH} = 3$. Yield per unit current varies in all cases between 89.6 and 99.7%, increasing with increasing current density and at high pH (3 to 4) values, and falling sharply at low current densities and at $\text{pH} = 1$ or 2 . In the next stage of the investigation the anodic processes were studied. Four types of electrolytes were used containing (in g/litre): (A) 25 $\text{Ni}(\text{BF}_4)_2$, 68.5 F_1 , 138.7 F_2 and 34.4 B; (B) same as (A) plus 15 $\text{NiCl}_2 \cdot 6\text{H}_2\text{O}$; (C) 240 $\text{NiSO}_4 \cdot 7\text{H}_2\text{O}$, 30 H_3BO_3 ; (D) same as (C) plus 20 NaCl . Cathodes were made of electrolytic nickel or of non-passivating nickel containing C 0.2%, Si 0.2% and S 0.005% (Ref 12). To reduce to minimum the difference between their true and the geometric surface areas, the cathodes were

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polished first mechanically and then electrolytically (to remove the plastically deformed layer). The results, in the form of graphs showing the relationship between the anodic current density and the anode potential, are reproduced in Figure 5: graph (1) - non-passivating nickel anode in electrolyte C; graph (2) - as (1) but electrolyte D used; graph (3) - non-passivating nickel anode in electrolyte A; graph (4) - as (3) but in electrolyte B; graphs (5) and (6) - electrolytic nickel anode in electrolytes C and D, respectively. When the ratio of the anode and cathode surface areas was $S_a/S_k = 2:1$, the yield of the dissolved metal per unit current was high even in the absence of chlorides in the electrolyte but for $S_a/S_k = 1:1$, a slightly lower yield was obtained. After the bath was operated for the equivalent of 800 A.hour per 1 litre of electrolyte, the concentration of the nickel-bearing salt in the solution was unchanged and only a slight rise in the pH number was observed. In their conclusions the authors state that: 1) the main advantage

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to be gained by using the fluoborate electrolyte instead of a sulphate solution for nickel plating is that the process can be intensified, i.e. higher current densities can be employed without affecting the quality of the deposit and yet without reducing the yield per unit current. The maximum permissible current density can be increased by increasing the Ni concentration in the electrolyte (up to 3N) and by using higher temperatures; ~~ii~~ when the concentration of H_3BO_3 in the electrolyte is increased, the cathode potential is reduced and so is the maximum permissible current density. In spite of this effect, which is probably due to partial dissociation of $Ni(BF_4)_2$ to $Ni(BF_3OH)_2$ and to a decrease in the activity of the nickel ions in the electrolyte, it is recommended to maintain the H_3BO_3 concentration slightly (approx.

15 g/litre) above the value corresponding to the stoichiometric ratio, since free H_3BO_3 improves the

Card6/8 stability of the electrolyte and makes it less reactive;

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Electrode Processes in Electro-deposition of Nickel from Fluoborate Electrolyte

iii) the effect of chlorides introduced in the electrolyte in the form of $\text{NiCl}_2 \cdot 6\text{H}_2\text{O}$ in amounts up to 15 g/litre is beneficial, since they reduce polarisation. However, at high (more than 30 g/litre) chlorides concentrations, the maximum permissible current density is lower and the quality of the deposit is adversely affected. At 50 °C, the effect of the chlorine ions becomes insignificant, most likely owing to the reduced adsorption of these ions on the cathode surface and to sufficiently high activity of the anode, particularly when made of non-passivating material. Owing to the latter factor, the fluoborate electrolytes are quite stable, so that in operation it is only necessary to replenish the fluoboric acid in order to maintain constant pH, since the anodic yield per unit current is slightly higher than the cathodic;

(iv) the optimum value of the pH number of a fluoborate solution for nickel plating is between 3 and 3.5. At pH = 1, the yield per unit current is too much affected by the variation of the current density, while at

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PANCHENKO, I. I., Cand of Tech Sci — (diss) "Electrolytic Nickel Plating by Cobalt and an Alloy of Nickel and Cobalt From Fluorinated Electrolytes," Moscow, 1959, 12 pp (Institute of Non-Ferrous Metallurgy im M. I. Kalinin) (KL, 4-60, 120)

SOV, 122-55-3-2, 34

AUTHORS: Layner, V.I., Doctor of Technical Sciences, Professor,
and Panchenko, I.I., Engineer

TITLE: Nickel Plating in Fluoroborate Electrolyte
(Nikelirovaniye vo ftorboratnom elektrolite)

PERIODICAL: Vestnik mashinostroyeniya, 1959, Nr 5, pp 65-68 (USSR)

ABSTRACT: An experimental study is reported concerned with the effect of each of the main factors on the nickel plating process in fluoroborate electrolytes. The solution of hydrofluoboric acid was prepared from hydrofluoric and boric acids by mixing the latter into the former in small doses accompanied by stirring and external cooling. Small doses of nickel carbonate were added accompanied by cooling. Concentrates with a specific gravity of 1.5 containing 180 g/litre of nickel can be obtained as fluoroborate of nickel. The fluoroborate of nickel has the formula $\text{Ni}(\text{BF}_4)_2 \cdot 6\text{H}_2\text{O}$. A method of analysis was used proposed by Ryss, I.G. ("The Chemistry of Fluorine and its Anorganic Compounds" Monograph, Goskhimizdat, 1956). Nickel was determined by the alkaline solution of di-methylglyoxime. 500 milli-litres of diluted

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Nickel Plating in Fluoroborate Electrolyte

concentrate were used in each test with a nickel anode and a cathode of copper, iron or brass foil. The current efficiency was measured with a copper coulometer. The specimens were electrochemically degreased, pickled in nitric acid (copper, brass) or hydrochloric acid (iron), washed in water then in alcohol, dried, cooled and weighed. Before immersing in the plating bath, the specimens were pickled in a 10% solution of sulphuric acid, washed in mains water and finally in distilled water. After plating, the specimens were washed in cold and hot water and in alcohol, dried, cooled and weighed. The curves of voltage against current density were plotted from measurements by the compensation method using a platinum disc cathode soldered into molybdenum glass. Before taking the readings, the cathode was covered by a 5 micron coat of copper from a sulphuric acid electrolyte. The deposited nickel was removed in hot 50% HNO_3 or by anodic dissolution in 50% H_2SO_4 .

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Nickel Plating in Fluoroborate Electrolyte

The anode was a disc of non-passivated nickel. Examining the effect of nickel salt concentration an increase from 1 N to 3 N solutions at a pH of 3.0 and 50°C increases the permissible current density. Deposits of 25 microns are obtained without burns and dendrite formation at the corners. Further increase of concentration is not profitable. The concentration of boric acid has no effect on the current efficiency and the quality of the deposit. Solutions with a small excess of boric acid (10 g/litre) are more stable. The effect of chlorine ion concentration was studied by using electrolytes without chlorine ions and those with 15, 30 and 50 g/litre $\text{NiCl}_2 \cdot 6\text{H}_2\text{O}$. Without chlorine ions, the quality of the deposit deteriorates, pitting increases, the current efficiency diminishes and the permissible current density drops. 15 g/litre is the best concentration. A temperature increase is beneficial by reducing polarization. The permissible current density increases and the quality of the deposit improves up to a temperature of 60°C. The pH value was

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Nickel Plating in Fluoroborate Electrolyte

studied in the range between 1 and 5. At a value of 4 and above the solution becomes unstable. At values of 1 and 2 the deposit contains much small pitting. The optimum value is 3. The buffer properties were studied at a temperature of 20°C by adding to fresh samples of 50 millilitre of electrolyte quantities of 10, 20 and 30 millilitre of 0.2N solution of NaOH and 0.2N solution of HCl and the measurement of the pH value of the solution by the quinhydrone method. The electrolytes differed in the nickel, boric acid and chloride content. Sulphuric acid electrolytes were examined alongside the fluoroborate electrolytes. It was shown that fluoroborate electrolytes have a substantially greater buffer property than sulphuric acid electrolytes with buffer admixtures. The stability of the electrolyte of the optimum composition, after plating under optimum conditions was judged by analysis for nickel, fluorine, boron and the pH value after 100 ampere-hours. A very good stability was observed. The anode to cathode surface ratio was varied between 8:1 and 1:1.

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Nickel Plating in Fluoroborate Electrolyte

After 300 ampere-hours per litre of electrolyte, the nickel, both forms of fluorine and boron contents did not change. The pH value rose from 3 to 3.4. The anode and cathode polarizations were plotted (Fig 6). The adhesion of deposits on iron, brass and copper specimens was judged by bending over 180° until fracture and by heating to 150°C. Deposits of 10, 30, 60, 100 and 500 microns were tested. Except in the thickest deposit, no delamination was found. In deposits beyond 30 microns, porosity was not observed. There are 6 figures, 1 table and 11 references, 8 of which are English, 2 Soviet and 1 French.

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PANCHENKO, Ivan Ivanovich; PROKOP'YEV, K.A., kand.tekhn.nauk, retsenzent;
SHEMONAYEV, A.S., inzh., red.; VASIL'YEVA, V.P., red.izd-va;
SHCHETININA, L.V., tekhn.red.

[Vibration resistance of turbine blades] Vibratsionnaya prochnost'
lopatok turbin. Moskva, Gos.nauchno-tekhn.izd-vo mashinostroit.
lit-ry, 1959. 253 p. (MIRA 12:9)
(Blades--Vibration)

VOROB'YEV, M.M.; IAN BIR, 197.

Distribution of 14 copies to the Ministry of Defense and Navy
Provinces; no. 33 no. 5:625
S-0 154. (MIRA 18:4)

PANCHENKO, I. P.

"Alloys of Titanium With Tungsten and Aluminum," by N. T. Gudotsov and I. P. Panchenko, Izvestiya Akademii Nauk SSSR, Otdeleniye Tekhnicheskikh Nauk, No 2, Feb 57, pp 139-143

An investigation of the hardness, endurance, and heat-resistance of alloys of titanium with 3 percent aluminum and 5, 10, and 15 percent tungsten revealed the expediency of alloying titanium with both aluminum and tungsten; these alloys are strengthened at 400-600 degrees [centigrade], as a result of which the hardness at room temperature is increased to 44, 82, and 85-117 Vickers units for the corresponding percentages; hardness and endurance increase with increased tungsten content. (U)

Sum. 1374

PPACHINIC 7 2
USSR/Microbiology - Antibiosis and Symbiosis, Antibiotics.

F-2

Abs Jour : Ref Zhur - Biol.: No 4, 1958, 14736

Author : Gromashevskaya, L.L., Golub, N.F., Panchenko, I.P.

Inst : -

Title : Sensitivity of Dysentery Bacteria to Biomycin.

Orig Pub : V sb.: Disenteriya, Kiev, Gosmedizdat USSR, 1956, 62-69

Abstract : Studies of the sensitivity of 120 strains of Flexner bacteria of various serotypes and 80 strains of Sonne bacteria to biomycin (I), levomycetin and norsulfazole showed that the most active I depressed the growth of the main mass of bacteria (91.5% of strains studied) at a concentration of 0.1-6.0 μ /ml. Sonne bacteria are more resistant to activity of I than Flexner bacteria, the sensitivity of which did not depend upon belonging to a definite serotype. The microorganisms are more sensitive to I on a Moore than on a Drobetko medium. I decreases oxygen consumption on a propifering as well as on a non-proliferating culture

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Card 2/2

Panchenko, I. P.

137-58-2-4253

Translation from: Referativnyy zhurnal. Metallurgiya, 1958, Nr 2, p 281 (USSR)

AUTHORS: Gudtsov, N. T., Panchenko, I. P.

TITLE: Titanium-tungsten Alloys (Titanovol'framovyye splavy)

PERIODICAL: Sb. Mosk. in-t stali, 1957, Vol 36, pp 5-12

ABSTRACT: A study was made of the effect of W on the microstructure, hardness, and long-term strength of magnesium-fusion-process Ti alloys (with 5, 10, and 15% W contents) smelted in graphite crucibles under a residual pressure of 10^{-2} mm Hg and containing 0.7-1.0% C. The alloys, quenched in water after being maintained 1-2 hours at a temperature of 1100°C , had the microstructure of a transformed β and molecular phase containing $\sim 10.5\%$ C. As an alloy's W content was increased, the microhardness of the molecular phase increased also, as a result of its enrichment with W. Quenching reduced the hardness of Ti-W alloys and increased their ρ . After being annealed at $\sim 800^{\circ}$ they had a three-phase structure: a molecular phase and two solid solutions based on α -Ti and W. The microstructure of alloys which had been aged at $200-500^{\circ}$ for 10-24 hours retained the acicular form it had acquired as a result of the quenching.

Card 1/2

137-58-2-4253

Titanium-tungsten Alloys

As the W content was increased, the formation temperature of the high-strength phase rose from 200° (in the case of the Ti) to 400-500° (in the case of the Ti/15%-W alloy). Gradual heating both intensified and raised the temperature of the dispersion-hardening process in the W-enriched alloys. Heating to 600° caused coagulation of the high-strength phase and a decline in hardness. At 650-700°, under a stress of 10-20 kg/mm², the investigated alloys exhibited a relatively low long-term strength, which increased, however, concurrently with their W content.

G. T.

1. Titanium alloys--Microstructure
2. Titanium alloys--Production
3. Titanium alloys--Test methods
4. Titanium alloys--Test results

Card 2/2

PANCHENKO, I.P.

Modernizing crank presses. Mashinostreitel' no.11:7-8 N '58.
(MIRA 11:12)

(Power presses)

PANCHENKO, I.P.; GUDTSOV, N.T. [deceased]

Investigation of titanium alloys with tungsten, aluminum,
beryllium and boron. Titan i ege splavy no. 1:11/123
'58. (MIRA 14:5)

1. Institut metallurgii AN SSSR.
(Titanium alloys—Metallography) (Creep of titanium—Testing)

PANCHENKO, I. P.

18(2)

PHASE II - ABSTRACTS

AB-1

Akademiya nauk SSSR. Institut metallurgii

Titan i ego splavy; metallurgiya i metallovedeniye (Titanium and its Alloys; Metallurgy and Physical Metallurgy) Moscow, Izd-vo AN SSSR, 1958. 209 p. 4,000 copies printed.

Resp. Ed.: N.V. Agayev, Corresponding Member, USSR Academy of Sciences; Ed. of Publishing House: V.B. Babushnikov; Tech. Ed.: A.A. Kiseleva.

INTRODUCTION: This book, of which a Phase I Exploitation (SOV/1200) has been prepared, is a collection of scientific papers devoted to the study of titanium and its alloys from three main points of view: physical metallurgy, forming, and welding. Special problems investigated include structural changes occurring during welding, determination of the content of harmful gases, development of industrial methods of rolling, and oxidation at various temperatures.

PART I. PHYSICAL METALLURGY

Card 1/43

Titanium and Its Alloys (Cont.)

AB-1

Gudtsov, N.T. and I.P. Panchenko (Institute of Metallurgy, USSR Academy of Sciences) Investigation of Titanium Alloys Containing Tungsten, Aluminum, Beryllium, and Boron 114

The aim of this investigation was to study the microstructure, hardness, and creep of the following alloys: 5 percent W + Ti; 5 percent W + 3 percent Al + Ti; 5 percent W + 3 percent Al + 0.1 percent Be + Ti; and 5 percent W + 3 percent Al + 0.1 percent B + Ti. The following materials were used for making the alloys: Mg-reduced titanium (99.93 percent), aluminum (99.99 percent), beryllium (99.3 percent), and ferroboration (20.45 percent B). The alloys were vacuum-melted in graphite crucibles. The ingots were forged at 800-1100° C into small rods, from which the test specimens were made. Quenching was carried out at 1100°. The specimens were heated in evacuated quartz tubes for 2 hours, after which the tubes were broken in water. Conclusions. 1) Alloying of titanium with W, Al, Be, and B reduced the creep rate 26-40 times at 500° C and under a tension of 5 kg/mm² during a 528-hour test period. 2) The alloys tested should be used at temperatures below 500°. 3) Measurement of hardness in a vacuum installation in the process of heating at progressively higher temperatures permits the determination of the hardening range,

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Titanium and Its Alloys (Cont.)

AB-1

and measurement of hardness during isothermal holding at heat makes it possible to trace the rate and duration of the hardening process and to make a proper choice of aging conditions for the alloys. There are 5 figures, 7 tables, and 3 references (2 Soviet and 1 English).

PART II. FORMING OF TITANIUM AND TITANIUM-BASE ALLOYS

Pavlov, I.M. (Institute of Metallurgy, USSR Academy of Sciences)
General Conditions for Forming Titanium and Its Alloys 124
Titanium and its alloys require special conditions for hot and cold forming. Cold deformation of Ti (alpha phase), as in other metals with a hexagonal structure (Mg, Zn, etc.), is accompanied by marked twinning. Cold ductility of Ti is greater than that of these other metals because of the greater number of possible slip and twinning planes. In a single cold-forming operation, Ti and its commoner alloys can be deformed by not more than 30 percent. With fractional (or repeated) deformation the figure can be brought up to 80 percent. Cold working in a particular direction causes anisotropy to develop. The crystal anisotropy is weak, but the mechanical anisotropy rises sharply when car-

Card 26/43

AUTHOR: Panchenko, I.P. SOV/117-58-11-5/36

TITLE: The Modernization of Crank Presses (Modernizatsiya krivoshipnykh pressov)

PERIODICAL: Mashinostroitel', 1958, Nr 11, pp 7 - 8 (USSR)

ABSTRACT: Crank presses of the firm "Berningauz", with a pressure of 180 tons and more, are used for cold drop forging. The friction clutch of these presses has been modernized by engineers of KhtZ. There is 1 diagram.

1. Forge presses--Performance 2. Clutches--Design

Card 1/1

PANCHENKO, I.P.

Mechanized conveying of carts. Mashinostroitel' no.11:23 № '60.
(MIRA 13:10)

(Conveying machinery)

BAKLANOV, Ivan Mikheylovich, kapitan-nastavnik; LANCHENKO, Ivan
Panteleyevich, kapitan-nastavnik; KOLCHAKOV, S., red.

[Sailing tankers to the Antarctic] Opyt plavaniia tankerov
v Antarktiku. Moskva, Transport, 1965. 74 p.
(114 184)

PANCHENKO, I. Ya.

"The Clinical Importance of Electrocardiographs in the Evaluation of the Condition of the Cardiovascular System in Cattle during Various Functional Disturbances." Cand Vet Sci, Moscow Technological Inst of the Meat and Dairy Industry, 6 Jan 55. (W, 27 sec 54)

Survey of Scientific and Technical dissertations presented at USSR Higher Educational Institutions (12)
SO: Sum. No. 556, 24 Jun55

PANCHENKO, I.Ya.; PUCHKOVA, S.M.

Condition of the organism of lambs poisoned with strontium-90.
Veterinariia 42 no.5:77-80 My '65. (MIRA 18:6)

1. 09111-67
ACC NRI AT6029427

(A)

SOURCE CODE: UR/0205/66/006/004/0625/0627

AUTHOR: Dobryakova, G. V.; Panchenko, I. Ya.; Povalyayov, A. P.

ORG: none

TITLE: Ratio of strontium to calcium upon passage from rations into dog skeletons

SOURCE: Radiobiologiya, v. 6, no. 4, 1966, 625-627

TOPIC TAGS: dog, isotope, calcium, strontium, biologic metabolism, food ration, bone

ABSTRACT: Tests were conducted for 180 days on 2 groups of dogs whose overall and mineral metabolism had been raised to a slightly positive calcium balance. The dogs were fed bone meal from animals who had received strontium⁹⁰ for a long time. Doses for groups 1 and 2 were 4.4×10^{-9} and 4.4×10^{-8} curie and 1 and 10 g calcium respectively. The animals were sacrificed after 30, 90 and 180 days and samples of rations, bone, blood, soft tissues, urine and feces were studied radiometrically and chemically. In group 2, the absolute strontium⁹⁰ accumulation in bone was only 2 fold that of group 1 although its content in the ration was 10 fold. Calcium and strontium absorption in group 2 was 1.5-2 fold that of group 1. No statistically valid difference was found for the calcium content of bones from the animals of groups 1 and 2. The same applied to soft tissues and blood. Results show that the uptake of calcium and strontium from the gastrointestinal tract and their accumulation in the

UDC: 578.088.91:612.014.482

Cord 1/2

L 09414-67

ACC NR: AP6029427

body occurs in a certain ratio; thus their metabolism in the body is apparently subject to the same rules. The uptake depends on the Ca/Sr ratio with a discriminatory correction for Sr. Orig. art. has: 2 tables.

SUB CODE: 06, 07/ SUBM DATE: 4Feb65/ ORIG REF: 002/ OTH REF: 004

Card 2/2

E. A.

C. A. Kryazev, June 1948

104. Capillary pyrometer for determining the specific gravity
of gasoline. (I. V. Pivneva and K. N. Ivanchenko (Nef. Akad.
1947, No. 2, 25-26; J. Ind. Petrol., 1948, 34, 302 a) — Tests with
an instrument made to the design of Lipkin (C. 1944, 94) showed
it to be easy to handle and to give results repeatable within
0.0002—0.0003 unit R. B. CLARK

463. CAPILLARY PYCNOMETER FOR DETERMINING SPECIFIC GRAVITY OF
GASOLINES. Plotneva, O. V. and Pangchenko, K. N.
(Neftyanoe Khoz., 1947, 25, (2), 58-60; Chem. Abstr.,
1947, 41, 6392).

PANCHENKO, Kh. N.

PA 411

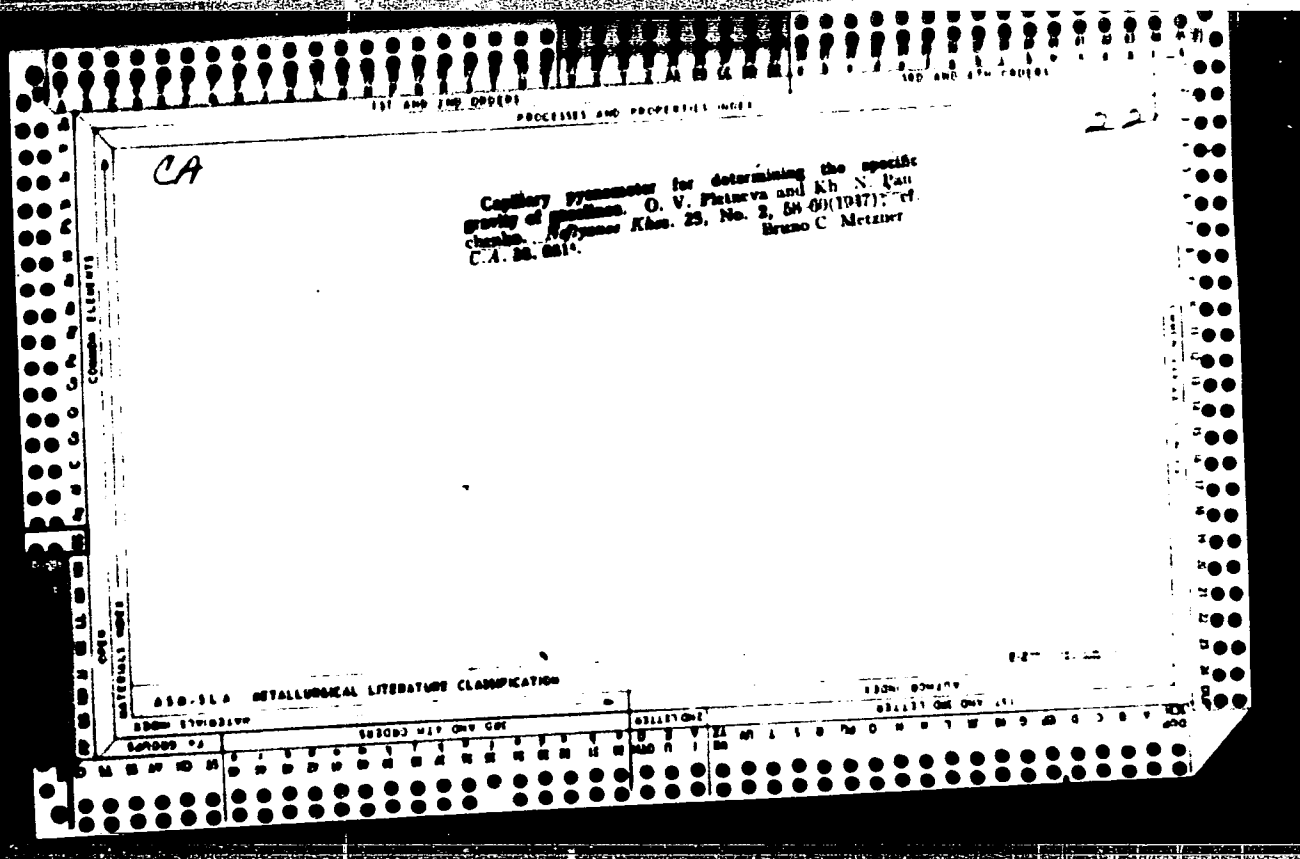
USSR/Chemistry - Fuels

Feb. 1977

"A Capillary Pycnometer to Determine the Specific Gravity of Gasoline,"
O. I. Pletneva, Kh. N. Panchenko, 2 pp

"Neftyanoye Khozjaysvo" Vol XXV, No 2

Schematic diagram and a graph. Discusses measurements of isopentane, heptane, benzol, isopropyl-benzol, and aviation gas, with a table.



1. PAMOTENKO, K.P., Engineer

2. KUP (COS)

(STI) (Stalinrad Tractor Plant)

"The Maximum Value of a Subsidiary Angle in a Star."
Stanki i Instrument, 12, No. 1, 1951

3. [REDACTED] Report K-1505, 4 Oct. 1951

PANCHENKO, K.P.

1952

Glasilo, A. N., Dobin, N. P., Zhevtunov, P. P., Naza-
rev, S. I., Panchenko, K. P., Popov, V. A., Popov, L. A.
and Stomov, M. V.: Tekhnologiya metallov (Technology
of Metals). Moscow: Mashiz. 1952. 637 pp. R. 21.
K. 10. Reviewed in Vestnik Mashinostroyeniya 34, No. 11,
100-8(1954).

of ⑦

GLADILIN, A. N.; DUBININ, N. P.; ZHVETUNOV, P.P.; NAZAROV, S.T.; PANCHENKO, K.P.;
POPOV, V.A.; POPOV, L.A.; and STOROSHEV, M.V.

Tekhnologiya Metallov, published by Mashgiz, Moscow, 1952

~~155x~~ Sum #148

PANCHENKO, F. P., ed.

Russian scientists - founders of the sciences of metalcutting. Moskv., Gos. Mashino-
tekhn. izd-vo mashinostroit. lit-ry, 1952. 27 p. (53-20056)

TJ1170.R95

ПА ПАНЧЕНКО, К. П.

123-1-604

Translation from: Referativnyy Zhurnal, Mashinostroyeniye, 1957,
Nr 1, p. 101 (USSR)

AUTHOR: Panchenko, K. P.

TITLE: Screw Profiled Form Cutters (Vintovyye fasonnyye rezbj)

PERIODICAL: Rezaniye metallov i instrument. Sbornik (MVTU, 54),
Moscow, Mashgiz, 1955, pp. 111-135

ABSTRACT: The author presents a mathematical analysis of the degree of precision of a screw type thread form cutter and a method of calculating design data. It was established that the most important factors affecting the cutting precision are the taper of ϕ_0 , the relief and rake angles, and the angle of profile distortion ϕ . Maximum error with $\phi_0 = 60^\circ$ and $\phi = 30^\circ$ is 1.4 mm. The longer the tapered part of the cutter (l) and the shorter the radius of the wider base (r), the greater the error. With $r = 50$ mm and $l = 30$ mm, the error reaches 1 to 1.2 mm. The effect of other factors is practically nil. On comparing the obtained data, the author concludes that screw profiled form cutters are as accurate as disc form cutters. B.L.Ya.

Card 1/1

ANTIPOV, K.F., inzhener; BILAKHIN, A.A., doktor tekhnicheskikh nauk,
 professor; BARYLOV, G.I., inzhener; BEYZSLIMAN, R.D., inzhener;
 BERDICHEVSKIY, Ye.G., inzhener; BOBKOV, A.A., inzhener; KALININ,
 M.A., kandidat tekhnicheskikh nauk; KOVAN, V.M., doktor tekhnicheskikh nauk,
 professor; KORINOV, V.S., doktor tekhnicheskikh nauk;
 KOSILOVA, A.G., kandidat tekhnicheskikh nauk; KUDRYAVTSEV, I.T.,
 doktor khimicheskikh nauk, professor; KURYSHEVA, Ye.S., inzhener;
 LAKHTIK, Yu.M., doktor tekhnicheskikh nauk, professor; MAYERMAN,
 M.S., inzhener; NOVIKOV, M.P., kandidat tekhnicheskikh nauk, PAKIY-
 SKIY, M.S., inzhener; PEREMOV, M.N., inzhener; POPILOV, A.Ye.,
 inzhener; POPOV, V.A., kandidat tekhnicheskikh nauk; SAVENIN, M.I.,
 doktor tekhnicheskikh nauk, professor; SASOV, V.V., kandidat tekhnicheskikh nauk,
 professor; SATAL, S.A., doktor tekhnicheskikh nauk, professor;
 SOKOLOVSKIY, A.P., doktor tekhnicheskikh nauk, professor; STANASVICH, V.G.,
 inzhener; PRUMIN, Yu.I., inzhener; KHRUKOV, V.I., inzhener;
 TSEYTLIN, L.B., inzhener; SHUKHOV, Yu.V., kandidat tekhnicheskikh nauk;
 BABIKIN, S.I., kandidat tekhnicheskikh nauk; GORODETSKIY, I.Ye.,
 VOLKOV, S.I., kandidat tekhnicheskikh nauk; GOROSHEIN, A.K., inzhener;
 DOSCHATOV, V.V., kandidat tekhnicheskikh nauk; ZAMALIN, V.I., inzhener;
 ISAYEV, A.I., doktor tekhnicheskikh nauk, professor; KUKHAROV, V.I.,
 kandidat tekhnicheskikh nauk; MALOV, A.N., kandidat tekhnicheskikh nauk;
 MARDANYAN, M.Ye., inzhener; PANCHENKO, K.P., kandidat tekhnicheskikh nauk;
 SEKRETEV, L.M., inzhener; STAYEV, K.P., kandidat tekhnicheskikh nauk;
 SYROVATCHENKO, P.V., inzhener; TAURIN, G.I., inzhener;
 AL'YASHOVA, M.A., kandidat tekhnicheskikh nauk;

(Continued on next card)